

Proposal for Human Respiratory Syncytial Virus Nomenclature below the Species Level

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Human respiratory syncytial virus (HRSV) is the leading viral cause of serious pediatric respiratory disease, and lifelong reinfections are common. Its 2 major subgroups, A and B, exhibit some antigenic variability, enabling HRSV to circulate annually. Globally, research has increased the number of HRSV genomic sequences available. To ensure accurate molecular epidemiology analyses, we propose a uniform nomenclature for HRSV-positive samples and isolates, and HRSV sequences, namely: HRSV/subgroup identifier/geographic identifier/unique sequence identifier/year of sampling. We also propose a template for submitting associated metadata. Universal nomenclature would help researchers retrieve and analyze sequence data to better understand the evolution of this virus.

Human respiratory syncytial virus (HRSV) is the leading cause of severe respiratory illness in children <5 years of age and is associated with substantial illness from lower respiratory tract infections in industrialized countries and substantial illness and death in low- and middle-income countries (1–5). HRSV also causes severe disease among elderly and high-risk adults (6).

In 2016, HRSV was reclassified by the International Committee on Virus Taxonomy (ICTV) into a new family, *Pneumoviridae*, genus, *Orthopneumovirus*, and species, *Human orthopneumovirus*. (7). The wider availability of viral sequencing technologies has increased submissions of HRSV sequences to databases (Figure 1), a trend we anticipate will continue.

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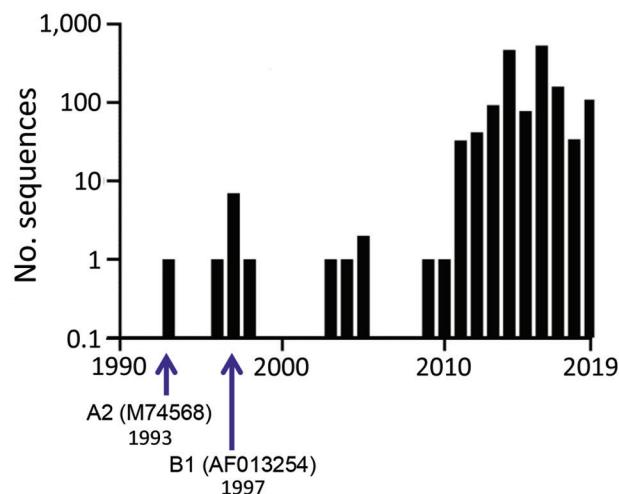


Figure 1. Annual numbers of HRSV whole-genome sequences released in GenBank since publication of the whole-genome sequence of HRSV A2, M74568, in 1993. HRSV, human respiratory syncytial virus.

Although ICTV provides nomenclature standards for virus taxa, there is currently no standardized format for HRSV nomenclature below the species level. Given the current interest in both HRSV and database submissions, a standard nomenclature is needed to simplify studies of the genomic diversity of HRSV strains and variants below the species level. ICTV's taxonomic reassignment provides us a timely opportunity to propose a universal naming convention for HRSV strains, sequences, and isolates, including a framework for database submissions that are rich in contextual information and associated metadata.

Several large laboratory HRSV surveillance and epidemiology studies are currently in progress. These studies include the World Health Organization's Global Respiratory Syncytial Virus (WHO RSV) Surveillance Project (<https://www.who.int/influenza/rsv>), which conducts large-scale testing for HRSV and extensive sequencing of HRSV-positive clinical specimens from >20 countries worldwide. Focused molecular analyses have helped elucidate HRSV household (8) and local (9) transmission dynamics and may guide development of strategies for the control of HRSV transmission. For example, molecular analysis showed that HRSV in health-care facilities can be acquired from sources within the facility or introduced from the community (10,11).

In temperate climates, annual HRSV epidemics usually occur in winter months; it remains to be seen how social distancing measures and nonpharmaceutical interventions due to the current coronavirus disease (COVID-19) pandemic will affect global HRSV circulation patterns. One of the 2 major genetic and antigenic HRSV subgroups, A or B, usually predominates in

alternating years, but both subgroups can also co-circulate in the same season. Early research has shown that subgroup A HRSV is associated with slightly greater clinical severity than subgroup B (12). Disease severity has been correlated with specific strains, genotypes, or clades, but to date, no consistent association has been established between strains (13–15), genotypes, or clades (16–19) and virulence. Thus, a possible role of different HRSV strains in disease severity remains to be elucidated. The lack of standard nomenclature and the scarcity of rich metadata in databases currently limit and complicate such studies.

Reliable and concise nomenclature systems below the species level are available for measles virus, influenza virus, rotavirus, filovirus isolates (20–23), and many other human viral pathogens. A similar nomenclature system tailored to HRSV and its pathology would support the requirements of researchers and the public health community by minimizing information errors when handling, storing, and shipping HRSV samples and when submitting, searching, and displaying sequencing data and associated metadata. Moreover, consistent nomenclature would improve the ability of researchers to pool and analyze data and associated information from different sources. To fill this need, an international group of researchers, in conjunction with the WHO RSV Global Surveillance Project, proposes a concise nomenclature system for HRSV below the species level.

HRSV Genome Organization

HRSV has a single-stranded nonsegmented negative-sense RNA genome ≈15,191–15,277 nt long (Figure 2, panel A) (7). The HRSV genome contains 10 genes, each encoding a separate mRNA with a single open reading frame (ORF) (Figure 2, panel A; Table), except for the M2 mRNA, which contains 2 overlapping ORFs. The 11 HRSV proteins are 2 nonstructural proteins (NS1 and NS2), nucleoprotein (N), phosphoprotein (P), matrix protein (M), small hydrophobic envelope protein (SH), attachment glycoprotein (G), fusion glycoprotein (F), the transcription processivity factor (M2-1), RNA regulatory factor (M2-2), and large RNA polymerase protein (L) (Table) (24,25). The F glycoprotein is the major viral neutralization and protective antigen, followed by the G glycoprotein (26).

HRSV Subgroups and Genotype Designations: Status and Outlook

HRSV subgroups A and B exhibit genomewide nucleotide and amino acid divergence (Figure 2, panel A) (25,27). The reference sequences for the 2 subgroups are derived from strains HRSV A2 (28–31;

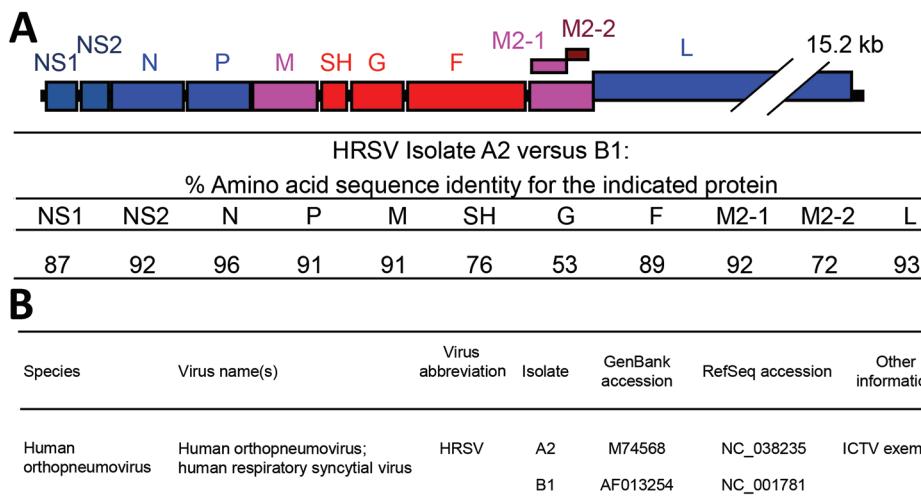


Figure 2. A) Schematic overview of the HRSV gene order and comparison of the amino acid identities of the reference strains of subgroups A (HRSV A2, GenBank accession no. M74568/NC_038235) and B (HRSV B1, GenBank accession no. AF013254/NC_001781). B) ICTV-proposed species designation, virus name, and associated GenBank reference sequences. HRSV, human respiratory syncytial virus; ICTV, International Committee on Virus Taxonomy; RefSeq, reference sequence.

GenBank accession number M74568.1; RefSeq accession number NC_038235) and HRSV B1 (32; GenBank accession number AF013254.1; RefSeq accession number NC_001781.1; Figure 2, panel B). F glycoprotein sequences between the 2 subgroups are well conserved (89% aa identity), whereas the G glycoproteins are the most divergent (53% aa identity between the subgroups) among the HRSV proteins (Figure 2, panel A) and undergo continuous molecular evolution. The ectodomain of the G glycoproteins of both subgroups contains a conserved central domain, representing an important antigenic site, flanked by 2 hypervariable domains (33). Except for the central conserved region, the antigenic cross-reactivity between G glycoproteins of the 2 subgroups is low (26).

Because the G ORF exhibits the greatest degree of genetic variability between isolates, it is most commonly used for studies on the molecular evolution

of HRSV. The genetic variability of HRSV strains over time has been commonly determined by sequencing the distal C-terminal third of the G ORF, which includes the second hypervariable domain. The variability in the G ORF is characterized by a high rate of nonsynonymous nucleotide changes, suggesting that evolution may be driven by immune pressure, even though this factor may be partially antibody independent (34). It is likely that variability in the G protein contributes to the capacity of HRSV to cause yearly outbreaks in the community (35–37). The nomenclature proposal outlined herein will be useful for the sequence analyses required to follow the molecular evolution of HRSV.

In a parallel effort, several research groups are working together on a genotyping proposal to provide a consensus on uniform genotype designations (38,39). As virus evolution continues, we expect new genotypes to emerge and older genotypes to become

Table. Widely accepted nomenclature for HRSV genes and proteins and gene annotation of strains HRSV A2 and HRSV B1*

Gene or genome region			Open reading frame genome position, nt			
Annotation†	Strain A2‡	Strain B1¶	Proteins		Strain A2	Strain B1
Leader region	1–44	1–44	Annotation	Abbreviation	Strain A2	Strain B1
NS1	45–576	45–577	Nonstructural protein 1	NS1	99–518	99–518
NS2	596–1098	594–1098	Nonstructural protein 2	NS2	628–1002	626–1000
N	1126–2328	1125–2327	Nucleoprotein	N	1141–2316	1140–2315
P	2330–3243	2331–3244	Phosphoprotein	P	2347–3072	2348–3073
M	3253–4210	3254–4208	Matrix protein	M	3262–4032	3263–4033
SH	4220–4629	4218–4630	Small hydrophobic protein	SH	4304–4498	4303–4500
G	4674–5596	4675–5600	Attachment glycoprotein	G	4689–5585	4690–5589
F	5649–7551	5653–7552	Fusion glycoprotein	F	5662–7386	7666–7390
M2	7598–8558	7609–8568	Matrix protein 2			
			Matrix protein M2–1	M2–1	7607–8191	7618–8205
			Matrix protein M2–2	M2–2	8160–8432	8171–8443
L	8491–15068	8501–15080	Polymerase protein	L	8499–14996	8509–15009
Trailer region	15069–15223	15081–15225				

*HRSV, human respiratory syncytial virus.

†Nucleotide annotations for leader and trailer region, and for indicated HRSV genes from the first nucleotide of the gene start signal [GGGGCAAAAT(A/G); GGAGCAAAT in case of the L gene] through the last adenine residue of the gene end signal [AGT(T/A)A(T/A/G)(A/T)(A/T)(A/T)A_n].

‡HRSV/A/USA/A2/2015 (45); GenBank accession number K992094.

¶HRSV/B/USA/B1/1985/B1 (46); GenBank accession number AF013254.1.



1. Virus name: human respiratory syncytial virus
2. HRSV subgroup identifier (A or B; X if unknown)
3. Country of sampling: ISO 3166-1 alpha-3 letter designation; XXX = unknown
4. Unique isolate identifier, restricted to 8 alphanumeric characters
5. Calendar year of sampling (YYYY) or XXXX if unknown

Figure 3. Schematic representation of the 5 consensus nomenclature elements of HRSV strains and isolates, with examples (top) and an explanation of each element (bottom). HRSV, human respiratory syncytial virus; ID, identification number; ISO, International Organization for Standardization.

extinct. HRSV genotyping designations will need to capture present molecular evolutionary status and be adaptable to changes and will need to be reevaluated periodically by a global consortium.

Nomenclature Proposal for HRSV Strains and Isolates

For molecular epidemiology studies, a concise standard for short identifiers of specific HRSV sequences, suitable for the short definition lines that give context to a sample and its derived sequence, would be useful. Ideally, concise standardized identifiers should convey key information about each individual sequence in an alignment or phylogram, including source, date, and type, if known. Here, we aim to define this type of common naming convention for HRSV samples and isolates. We also propose the use using standard names and appropriate annotations for HRSV genes, provide examples to guide the annotation of sequence data during the sequence submission process, and suggest how to submit metadata associated with the source materials of HRSV sequences.

Definition Lines for HRSV Sequence Submissions

GenBank records available through the National Center for Biotechnology Information (NCBI) are identified by 2 elements: a unique alphanumeric accession number and a definition line. The definition line is the portion of the identifier commonly associated with GenBank records shown in BLAST results and other searches. Definition lines are generated by the submitter during the sequence submission process and include the species and isolate name (e.g., *Human orthopneumovirus* isolate HRSV/A/USA/001/2011, complete genome [proposed]). (40). We propose a standardized format to capture 5 sample-specific parameters of HRSV-positive clinical samples or iso-

lates to be included in sequence definition lines compatible with database naming requirements (Figure 3), in this specific order: [virus name abbreviation]/[HRSV subgroup]/[geographic identifier]/[unique sequence identifier]/[year of sampling].

Elements of Sequence Definition

I. Organism name; virus name abbreviation: HRSV

ICTV's species name, *Human orthopneumovirus* (7), will be reflected as HRSV in the NCBI definition line. During submission to databases, the organism name can be entered as either Human orthopneumovirus or human respiratory syncytial virus. The abbreviation HRSV should be used in the definition line regardless of which organism name is provided.

II. HRSV subgroup: A or B; X, if unknown

III. Geographic identifier for the location of sampling.

Because individual HRSV research networks have predefined requirements, we suggest some flexibility for this field:

- i. If not specified by a research network, the ISO 3166-1 α-3 letter country code (<https://www.iso.org/iso-3166-country-codes.html>) should be used to indicate the country of sampling (XXX, if unknown). We strongly suggest that submitters provide any more specific geographic information on the sampling location (e.g., city or state) in metadata fields rather than in the definition line.
- ii. The WHO Global RSV Surveillance Project plans to use just the simple English-language name for the country.
- iii. Individual national studies may require a state, province, or city designation, in addition to the mandatory country name. If required, a period should be used to set off the country name ([state/province/city].[country name]).

IV. Unique isolate identifier

This field must be restricted to 8 alphanumeric characters. Underscores are permitted, but neither other special characters (e.g., /, %, \$, @) nor spaces can be used. Controversial names or phrases, names of prominent people, and trademarked names or phrases cannot be used. To prevent duplication of sample or isolate identifiers by different groups, we recommend inserting a letter code identifying a study or institute before the sample number. For example, unique isolate identifiers for samples from the INFORM-RSV study might use "INF" followed by a number (e.g., INF001 in HRSV/A/COUNTRY/INF001/2019).

V. Year of sampling; YYYY or XXXX, if unknown.**Examples of Sequence Definition Lines Using Proposed Nomenclature**

- HRSV/A/USA/001/2011
- HRSV/B/Denver.USA/14617/1985
- HRSV/A/IRN/001/2017
- HRSV/A/Iran/001/2017
- HRSV/X/IRN/001/2017 (subgroup unknown)
- HRSV/B/New_Zealand/FR123/2020

Our nomenclature proposal prioritizes a short, concise definition line that will be easy to use in the laboratory, easily readable, and be a uniform system for HRSV in public databases. Additional host, virus, location or temporal information if desired could be submitted in metadata fields, which would allow researchers, epidemiologists, and database users to apply specific metadata filters, as needed for data retrieval and specific applications, analyses, or for displaying designations, such as in dendrograms.

Terminology for Annotations

To support efficient data analysis, uniform designations must be used at the database submission stage. Commonly accepted names for HRSV genes and proteins are shown in the table. An HRSV gene comprises a gene start signal GGGGCAAAT(A/G), an ORF with adjacent noncoding regions, if present, and the gene end signal through the last adenosine residue [AGT(T/A)A(T/A/G)(A/T)(A/T)(A/T)A_n] (Figure 4; 41). Each HRSV gene contains a single ORF, except for the M2 gene, which has 2 overlapping ORFs, M2-1 and M2-2. Nucleotide annotations of genes and ORFs for the HRSV A2 (Figure 4) and HRSV B1 reference sequences are shown in the table.

Metadata for HRSV Sequence Submissions

What is the most pertinent host data will depend on the interests and objectives of individual study groups. For example, when studying HRSV in a pediatric setting, prematurity may be of interest, but when studying HRSV in an adult setting, researchers may be more interested in whether participants are immunocompromised. We suggest information that could be included in metadata fields for HRSV:

**1. Isolation source: sample type
(upper or lower airways)**

Viral RNA can be extracted directly from a clinical sample, from an isolate grown in cell culture, or possibly from a cDNA-derived recombinant virus. The sources of sequences from isolated viruses can be

identified by the following designations:

- wt: wild-type; sequences derived from RNA extracted directly from clinical specimens
- tc: tissue culture; sequences derived from RNA extracted from HRSV isolates propagated in tissue culture
- rec: recombinant; sequences of cDNA-derived recombinant virus (including vaccine strain)

2. Host

Homo sapiens; subject age. Indicate years and months if <5 years of age, years only if ≥5 years of age. Sex should be spelled out if known.

3. Country, state, and (nearest) city of sampling

Metadata information must include the full country name (not the 3-letter abbreviation) from the NCBI list of accepted country designations (<https://www.ncbi.nlm.nih.gov/GenBank/collab/country>). City, state, or province can also be included. Names should be written based on the standard ASCII letters including spaces if required (https://www.nist.gov/system/files/documents/2021/03/23/ansi-nist_2010_traditional_|encoding.pdf). Geolocation coordinates of the location where sampling took place should be included if known.

4. Collection date

We highly recommend that the exact date of specimen collection (DD-Mon-YYYY format; e.g., 17-Feb-2002) be used; if exact date is not known, at least the month and year should be indicated (Mon-YYYY format).

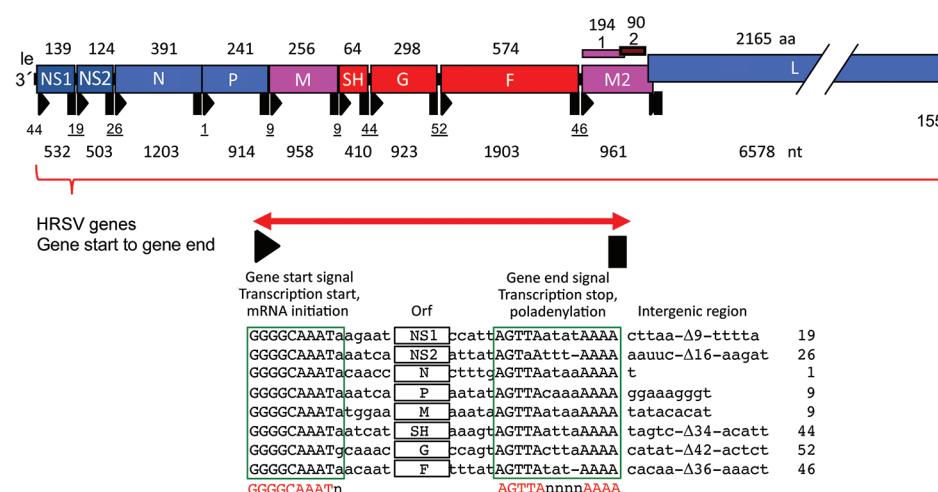
5. Genotype according to the consensus in genotype classification by an HRSV working group [in progress (38,39)]

Associated with the International RSV Society, a special interest group of the International Society for Influenza and other respiratory viruses <https://www.isirv.org/site/index.php/special-interest-groups/international-respiratory-syncytial-virus-society>.

6. Metadata on the patient-host and the clinical disease should be included in the notes field in a structured format

Protected personally identifiable health information will be excluded from metadata submissions.

- If >6 months of age, birthweight and gestational age at birth.
- Significant pediatric co-morbidities, including prematurity, congenital cardiac disease, and broncho-pulmonary dysplasia (BPD).
- Twin? (Y/N)



- Exposed to specific HRSV therapeutic, vaccine, antibody, or antiviral? (Y/N)
- Viral or bacterial co-infections, if known; pathogen species should be spelled out.
- Adult underlying conditions, such as chronic obstructive pulmonary disease (COPD) or asthma, or altered immune status (e.g., immunocompromised, bone marrow transplant recipient).
- Disease outcomes. Five grades are distinguished:
 - No medical care.
 - Outpatient or emergency room.
 - Hospital admission.
 - ICU admission.
 - Death.

For NCBI submissions, data can be entered through the web interface, or uploaded as tab-delimited text files. Sequences can be uploaded in FASTA format (<https://blast.ncbi.nlm.nih.gov>; Appendix, <https://wwwnc.cdc.gov/EID/article/27/6/20-4608-App1.pdf>), with associated metadata provided in a plain text, tab-delimited, source modifier table (Appendix Table 1) and gene or protein annotations provided in a plain text, tab-delimited, 5-column feature table (Appendix Table 2).

Outlook

Molecular surveillance has revealed that multiple HRSV genotypes circulate simultaneously in communities. Circulating genotypes often vary between communities, and circulation patterns within a community can change from year to year. Extended monitoring of circulating viruses is necessary to better understand transmission and molecular evolution (42). As HRSV vaccine candidates and antivirals are being developed, molecular epidemiology studies may reveal potential

effects of prevention strategies on viral evolution and possible antibody-escape variants. Timely sharing of HRSV data worldwide through the use of public databases is essential. We propose that sequence data be uploaded to publicly accessible databases, such as NCBI (31). Although NCBI is the most complete repository for HRSV sequence information, studies may require that sequences first be submitted to other databases, such as GISAID (<https://www.gisaid.org>).

Public access will provide useful availability for investigators to submit, query, and analyze HRSV sequence data, enabling the evolutionary analysis of sequence diversity within or between HRSV genotypes. The utility of public access has been clearly demonstrated with the emergence of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) and the critical role that genetic sequence analysis has played. Notably, a nomenclature model for SARS-CoV-2 similar to what we have proposed for HRSV has been adopted, although some differences remain between databases (e.g., NCBI, SARS-CoV-2/human/USA/COVID20-1096/2020; GISAID, hCoV-19/Australia/VIC12/2020). When an HRSV vaccine becomes available, high-quality, geographically representative, country-specific data on circulating strains and rich datasets of well-curated, standardized, and parsable data will be required to monitor and trace possible evolutionary changes in response to vaccine-induced selective pressure (43,44). This proposal will profit from strong support by members of the International RSV Society, a special interest group of the International Society for Influenza and other Respiratory Virus Diseases (<https://isirv.org>), members of the WHO Global RSV Surveillance Project, and the HRSV research community.

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S.H., S.J., and W.Z. work with the World Health Organization. The authors are responsible for the views expressed in this publication and they do not necessarily represent the decisions, policy or views of the World Health Organization. Names of specific vendors, manufacturers, or products are included for public health and informational purposes; inclusion does not imply endorsement of the vendors, manufacturers, or products by the World Health Organization. L.A. has done paid consultancies on RSV vaccines for Bavarian Nordic, Novavax, ClearPath Vaccines Company, and Pfizer; his laboratory is currently receiving funding through Emory University from Pfizer and Advac for laboratory studies for HRSV surveillance studies in adults, and he holds a subcontract on an NIH SBIR award to Sciogen on G protein HRSV vaccines. L.A. is a co-inventor on several CDC patents on the HRSV G protein and its CX3C chemokine motif relative to immune therapy and vaccine development, and on a patent filing for use of HRSV platform VLPs with the F and G proteins for vaccines. U.B. reports CRADA support to NIH from Sanofi Pasteur, outside the submitted work; in addition, she has patents on live-attenuated HRSV with royalties paid to NIH by Sanofi Pasteur. There are no additional conflicts of interest by any of the authors.

About the Author

Dr. Salimi is a virologist, associate professor, and head of the virology department at the School of Public Health, Tehran University of Medical Sciences. His primary research interests include genetic characterization, immunopathogenesis, and vaccine design of respiratory viruses.

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May 2021

COVID-19

- Coordinated Strategy for a Modeling-Based Decision Support Tool for COVID-19, Utah, USA
- Clinical Laboratory Perspective on Human Infections Caused by Unusual Nonhemolytic, Lancefield Group B *Streptococcus halichoeris*
- Case Series of Laboratory-Associated Zika Virus Disease, United States, 2016–2019
- Successful Control of an Onboard COVID-19 Outbreak Using the Cruise Ship as a Quarantine Facility, Western Australia T
- Coccidioidomycosis and COVID-19 Co-Infection, United States, 2020
- Epidemiologic Findings From Case Investigations and Contact Tracing of the First 200 Cases of Coronavirus Disease 2019 (COVID-19) Identified in Santa Clara County, California, USA
- SARS-CoV-2 in Nursing Homes after 3 Months of Serial, Facility-Wide Point Prevalence Testing, Connecticut, USA
- Transmission of Severe Acute Respiratory Syndrome Coronavirus 2 during Border Quarantine and Air Travel, New Zealand (Aotearoa)



- Engineered NS1 Provides Sensitive and Specific Zika Diagnosis from Patient Serology
- Characteristics and Clinical Implications of Carbapenemase-Producing *Klebsiella pneumoniae* Colonization and Infection, Italy
- Serologic Screening of Severe Acute Respiratory Syndrome Coronavirus 2 Infection in Cats and Dogs during First Coronavirus Disease Wave, the Netherlands
- Epidemiologic History and Genetic Diversity Origins of Chikungunya and Dengue Viruses, Paraguay
- Symptom Diary-Based Analysis of COVID-19 Disease Course, Germany, 2020
- Herd Immunity against Severe Acute Respiratory Syndrome Coronavirus 2 Infection in 10 Communities, Qatar
- Monitoring SARS-CoV-2 Circulation and Diversity through Community Wastewater Sequencing, the Netherlands and Belgium
- Active Case Finding of Current Bornavirus Infections in Human Encephalitis Cases of Unknown Etiology, Germany, 2018–2020
- Susceptibility to SARS-CoV-2 of Cell Lines and Substrates Commonly used to Diagnose and Isolate Influenza and Other Viruses
- Global Trends in Norovirus Genotype Distribution among Children with Acute Gastroenteritis
- Genetic Evidence and Host Immune Response in Persons Reinfected with SARS-CoV-2, Brazil
- Epidemiology of Confirmed COVID-19 Deaths in Adults, England, March–December 2020
- Prescribing Antimicrobial Drugs for Acute Gastroenteritis, Primary Care, Australia, 2013–2018

Proposal for Human Respiratory Syncytial Virus Nomenclature below the Species Level

Appendix

Appendix Table 1. Virtual examples of typical sequence-associated data, not based on existing primary data*

Seq	Isolate	Source	Sex/ age	Lab host	Location	Date	Geno	Notes	Sub
rsv 011	HRSV/A/PHL/pim16223/2016	Wt; sputum	M/ 3 mo	NA	Philippines	Dec 2016	ON1	Wt/gest age at birth: 7 lb/38 w; twin: no; HRSV therapy: no; co-infection: Strep A; severity: ER	A
rsv 012	HRSV/B/ARG/352162/2019	Wt; nasopharyngeal aspirate	F/ 2 y, 7 mo	NA	Buenos Aires, Argentina	May 2019	TBD	Co-morbidities: broncho-pulmonary dysplasia; severity: hospital admission	B
rsv 013	HRSV/A/Piura.PER/PIU048/2020	TC; nasal swab	F/ 87 y	HEp-2 cells x2	Piura, Peru	Apr 2020	TBD	Co-infection: influenza A; severity: no medical care	A

*Geno, genotype; lb, pound; Seq, sequence identification; Sub, subtype; TBD, to be determined; TC, tissue culture; wt, weight

Appendix Table 2. Virtual examples of typical sequence-associated data, not based on existing primary data*

Start location (first nt)	Stop location (last nt)	Feature name	Qualifier (product or gene)	Qualifier value (designation)
>Feature rsv011				
4	576	Gene	Gene	NS1
99	518	CDS	Product	Nonstructural protein 1
596	1098	Gene	Gene	NS2
628	1002	CDS	Product	Nonstructural protein 2
112	2326	Gene	Gene	N
1140	2315	CDS	Product	Nucleoprotein
2330	3242	Gene	Gene	P
2347	3072	CDS	Product	Phosphoprotein
3246	4203	Gene	Gene	M
3255	4025	CDS	Product	Matrix protein
4211	4621	Gene	Gene	SH
4295	4489	CDS	Product	Small hydrophobic protein
4666	5659	Gene	Gene	G
4681	5646	CDS	Product	Attachment glycoprotein
5713	7615	Gene	Gene	F
5726	7450	CDS	Product	Fusion glycoprotein
7660	8620	Gene		

Start location (first nt)	Stop location (last nt)	Feature name	Qualifier (product or gene)	Qualifier value (designation)
7669	8253	CDS	Gene	M2
8222	8494	CDS	Product	M2-1 protein
8552	15124	Gene	Product	M2-2 protein
8561	15058	CDS	Gene	L
			Product	Polymerase protein
>Feature rsv012				
4	576	Gene	Gene	NS1
99	518	CDS	Product	Nonstructural protein 1
596	1098	Gene	Gene	NS2
628	1002	CDS	Product	Nonstructural protein 2
112	2326	Gene	Gene	N
1140	2315	CDS	Product	Nucleoprotein
2330	3242	Gene	Gene	P
2347	3072	CDS	Product	Phosphoprotein
3246	4203	Gene	Gene	M
3255	4025	CDS	Product	Matrix protein
4211	4621	Gene	Gene	SH
4295	4489	CDS	Product	Small hydrophobic protein
4666	5659	Gene	Gene	G
4681	5646	CDS	Product	Attachment glycoprotein
5713	7615	Gene	Gene	F
5726	7450	CDS	Product	Fusion glycoprotein
7660	8620	Gene	Gene	M2
7669	8253	CDS	Product	M2-1 protein
8222	8494	CDS	Product	M2-2 protein
8552	15124	Gene	Gene	L
8560	15058	CDS	Product	Polymerase protein
>Feature rsv013				
<1	333	Gene	Gene	NS1
<1	275	CDS	Product	Nonstructural protein 1
353	855	Gene	Gene	NS2
385	759	CDS	Product	Nonstructural protein 2
882	2085	Gene	Gene	N
897	2072	CDS	Product	Nucleoprotein
2095	3007	Gene	Gene	P

Start location (first nt)	Stop location (last nt)	Feature name	Qualifier (product or gene)	Qualifier value (designation)
2112	2837	CDS	Product	Phosphoprotein
3011	3968	Gene	Gene	M
3020	3790	CDS	Product	Matrix protein
3976	4385	Gene	Gene	SH
4060	4254	CDS	Product	Small hydrophobic protein
4430	5351	Gene	Gene	G
4445	5338	CDS	Product	Attachment glycoprotein
5405	7307	Gene	Gene	F
5418	7142	CDS	Product	Fusion glycoprotein
7352	8312	Gene	Gene	M2
7361	7945	CDS	Product	M2-1 protein
7914	8186	CDS	Product	M2-2 protein
8245	>14615	Gene	Gene	L
8253	>14615	CDS	Product	Polymerase protein

*CDS, coding sequence (coding region of gene)

Human respiratory syncytial virus genome

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>rsv011
ACCGCAAAAATGCGTACAACAACTTGCATAACCAAAAAATGGGCAAATAAGAATTGATAAGTACCACTTAAATT
TAACTCCTTGGTTAGAGATGGGAGCAACTCATTGAGTATGATAAAAGTTAGATTGCAAATCTGTTGACAATGATGA
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```

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